



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) Publication number:

0 446 846 A2

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 91103709.1

(51) Int. Cl. 5: B41M 5/38

(22) Date of filing: 11.03.91

(30) Priority: 13.03.90 US 493076

(71) Applicant: EASTMAN KODAK COMPANY
343 State Street
Rochester, NY 14650-2201(US)

(43) Date of publication of application:
18.09.91 Bulletin 91/38

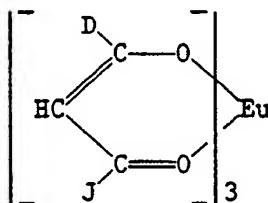
(72) Inventor: Byers, Gary Wayne, c/o EASTMAN
KODAK COMPANY
Patent Department 343 State Street
Rochester New York 14650-2201(US)
Inventor: Chapman, Derek David, c/o
EASTMAN KODAK COMPANY
Patent Department 343 State Street
Rochester New York 14650-2201(US)

(84) Designated Contracting States:
AT BE CH DE DK ES FR GB GR IT LI LU NL SE

(74) Representative: Brandes, Jürgen, Dr.Rer.Nat.
et al
Wuesthoff & Wuesthoff, Patent- und
Rechtsanwälte, Schweigerstrasse 2
W-8000 München 90(DE)

(54) Thermally-transferable fluorescent europium complexes.

(57) A donor element for thermal transfer comprising a support having on one side thereof a fluorescent europium complex dispersed in a polymeric binder. In a preferred embodiment, the complex has the formula:



wherein:

D is a substituted or unsubstituted, aromatic, 5- or 6-membered carbocyclic or heterocyclic moiety; and
J is -CF₃, -CH₃, -CH₂F or -CHF₂.

In another preferred embodiment, the europium atom may also have attached thereto at least one monodentate ligand with an electron-donating oxygen or nitrogen atom or at least one bidentate ligand with two electron-donating oxygen, nitrogen or sulfur atoms capable of forming a 5- or 6-membered ring with the europium atom.

EP 0 446 846 A2

This invention relates to fluorescent donor elements used in thermal transfer.

In recent years, thermal transfer systems have been developed to obtain prints from pictures which have been generated electronically from a color video camera. According to one way of obtaining such prints, an electronic picture is first subjected to color separation by color filters. The respective color-separated images are then converted into electrical signals. These signals are then operated on to produce cyan, magenta and yellow electrical signals. These signals are then transmitted to a thermal printer. To obtain the print, a cyan, magenta or yellow dye-donor element is placed face-to-face with a dye-receiving element. The two are then inserted between a thermal printing head and a platen roller. A line-type thermal printing head is used to apply heat from the back of the dye-donor sheet. The thermal printing head has many heating elements and is heated up sequentially in response to the cyan, magenta and yellow signals. The process is then repeated for the other two colors. A color hard copy is thus obtained which corresponds to the original picture viewed on a screen. Further details of this process and an apparatus for carrying it out are contained in U.S. Patent No. 4,621,271 by Brownstein entitled "Apparatus and Method For Controlling A Thermal Printer Apparatus," issued November 4, 1986.

The system described above has been used to obtain visible dye images. However, for security purposes, to inhibit forgeries or duplication, or to encode confidential information, it would be advantageous to create non-visual ultraviolet absorbing images that fluoresce with visible emission when illuminated with ultraviolet light.

U.S. Patent 4,627,997 discloses a fluorescent thermal transfer recording medium comprising a thermally-meltable, wax ink layer. In that system, the fluorescent material is transferred along with the wax material when it is melted. Wax transfer systems, however, are incapable of providing a continuous tone. Further, the fluorescent materials of that reference are incapable of diffusing by themselves in the absence of the wax matrix. It is an object of this invention to provide fluorescent materials useful in a continuous tone system which have sufficient vapor pressure to transfer or diffuse by themselves from a donor element to a dye-receiver.

U.S. Patents 4,876,237, 4,871,714, 4,876,234, 4,866,025, 4,860,027, 4,891,351, and 4,891,352 all relate to thermally-transferable fluorescent materials used in a continuous tone system. However, none of those materials fluoresce a visible red color when illuminated with ultraviolet light. A red color is desirable for many security applications. It is an object of this invention to provide a red fluorescent material for thermal dye transfer.

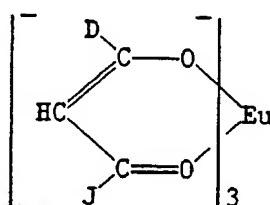
These and other objects are achieved in accordance with this invention which comprises a donor element for thermal transfer comprising a support having on one side thereof a fluorescent europium complex dispersed in a polymeric binder.

In a preferred embodiment of the invention, the europium complex has the formula:

35

40

45



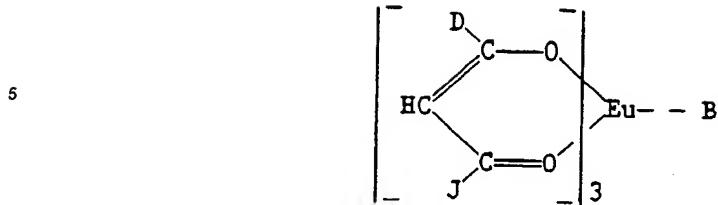
wherein:

D is a substituted or unsubstituted, aromatic, 5- or 6-membered carbocyclic or heterocyclic moiety, e.g., phenyl, 2-thienyl, 2-furyl, 3-pyridyl, etc.; and

50 J is -CF₃, -CH₃, -CH₂F or -CHF₂.

In another preferred embodiment of the invention, the europium atom may also have one or more "auxiliary" monodentate or bidentate ligands attached thereto which substantially increases the fluorescence. Thus, these "higher" coordinate complexes have the following formula:

55



wherein:

D and J are defined as above and B represents at least one monodentate ligand with an electron-donating oxygen or nitrogen atom, e.g., tri-n-octylphosphine oxide, pyridine-N-oxide or triphenylphosphine oxide; or at least one bidentate ligand with two electron-donating oxygen, nitrogen or sulfur atoms capable of forming a 5-or 6-membered ring with the europium atom, e.g., 2,2'-bipyridine, 1,10-phenanthroline, ethylene diamine or 1,2-diaminobutane.

The above fluorescent europium complexes are essentially non-visible, but emit with a unique red hue in the region of 610 to 625 nm when irradiated with 360 nm ultraviolet light. This red hue is highly desirable for security-badging applications.

20 Europium(III) is the only rare-earth known to be suitable for the practice of the invention. Rare earth metals, including europium, are described in the literature such as S. Nakamura and N. Suzuki, Polyhedron, 5, 1805 (1986); T. Taketatsu, Talanta, 29, 397 (1982); and H. Brittain, J.C.S. Dalton, 1187 (1979).

Compounds included within the scope of the invention include the following europium complexes derived from three beta-diketone ligands and optionally an auxiliary uncharged ligand(s):

25

30

35

40

45

50

55

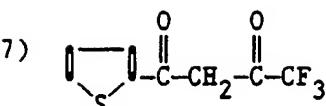
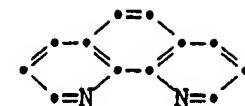
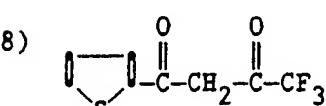
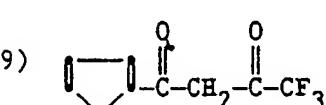
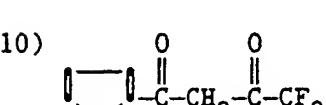
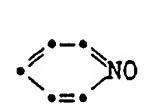
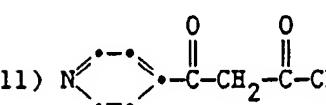
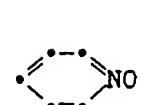
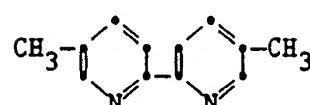
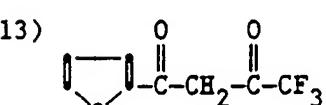
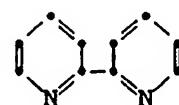
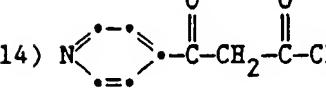
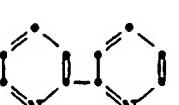
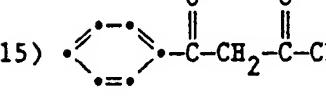
	A. Diketone Ligand	B. Auxiliary Ligand	Ratio Eu:A:B
5 1)		None	1:3:0
10 2)		None	1:3:0
15 3)		None	1:3:0
20 4)		None	1:3:0
25 5)		None	1:3:0
30 35 6)			1:3:1

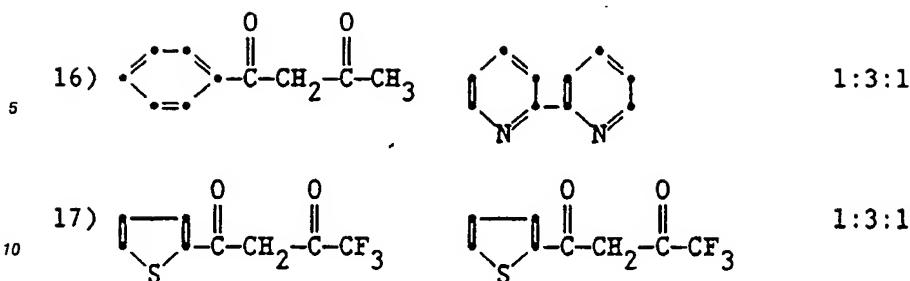
40

45

50

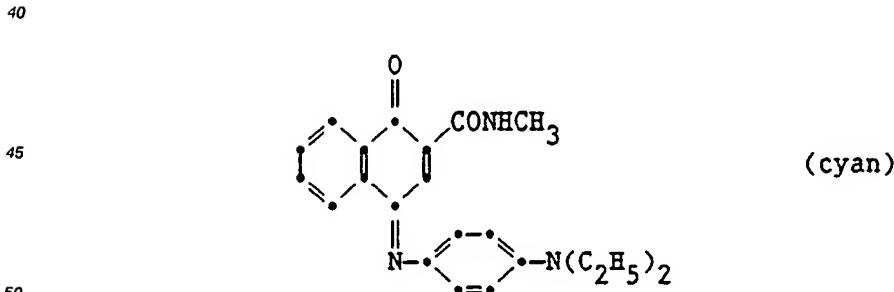
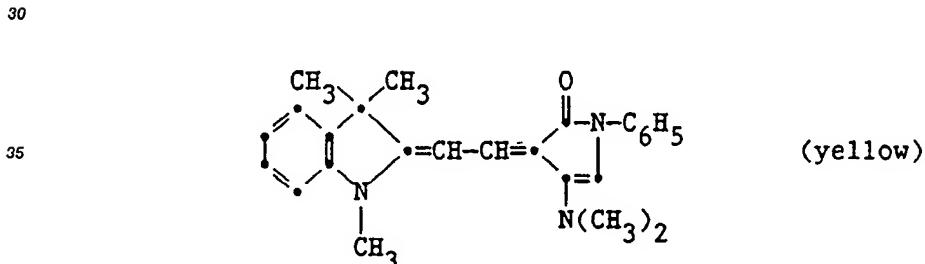
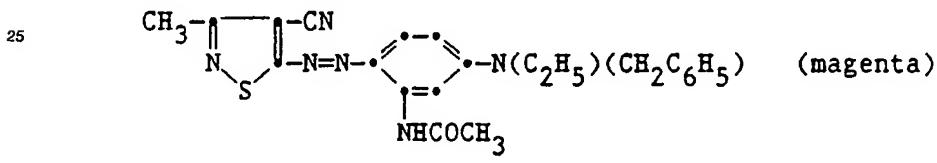
55

5	7) 		1:3:1
10	8) 	H ₂ NCH ₂ CH ₂ NH ₂	1:3:1
15	9) 	(C ₆ H ₅) ₃ PO	1:3:2
20	10) 		1:3:1
25	11) 		1:3:1
30	12) 		1:3:1
35	13) 		1:3:1
40	14) 		1:3:1
45	15) 	(n-C ₈ H ₁₇) ₃ PO	1:3:2



15 *This compound differs from the others in that it has four thenoyltrifluoroacetones coordinated with one europium, thus is octacoordinate and charged. To provide a neutral molecule, a salt is formed with tetramethylammonium as a counter-ion. It was found to have sufficient solubility to be coated and evaluated.

20 A visible dye can also be used in a separate or the same area of the donor element of the invention provided it is transferable to the dye-receiving layer by the action of heat. Especially good results have been obtained with sublimable dyes, such as:



50 or any of the dyes disclosed in U.S. Patent 4,541,830. The above dyes may be employed singly or in combination to obtain a monochrome. The above image dyes and fluorescent dye may be used at a coverage of from 0.01 to 1 g/m², preferably 0.1 to 0.5 g/m².

55 The fluorescent material in the donor element of the invention is dispersed in a polymeric binder such as a cellulose derivative, e.g., cellulose acetate hydrogen phthalate, cellulose acetate, cellulose acetate propionate, cellulose acetate butyrate, cellulose triacetate; a polycarbonate; poly(styrene-co-acrylonitrile), a poly(sulfone) or a poly(phenylene oxide). The binder may be used at a coverage of from 0.1 to 5 g/m².

The fluorescent material layer of the donor element may be coated on the support or printed thereon by a printing technique such as a gravure process.

Any material can be used as the support for the donor element of the invention provided it is dimensionally stable and can withstand the heat of the thermal printing heads. Such materials include 5 polyesters such as poly(ethylene terephthalate); polyamides; polycarbonates; glassine paper; condenser paper; cellulose esters such as cellulose acetate; fluorine polymers such as polyvinylidene fluoride or poly(tetrafluoroethylene-co-hexafluoropropylene); polyethers such as polyoxymethylene; polyacetals; polyolefins such as polystyrene, polyethylene, polypropylene or methylpentane polymers; and polyimides such as 10 polyimide-amides and polyether-imides. The support generally has a thickness of from 2 to 30 μm . It may also be coated with a subbing layer, if desired.

When using the donor element of the invention with a resistive head, the reverse side of the donor element is coated with a slipping layer to prevent the printing head from sticking to the donor element. Such a slipping layer would comprise a lubricating material such as a surface active agent, a liquid lubricant, a solid lubricant or mixtures thereof, with or without a polymeric binder.

15 The receiving element that is used with the donor element of the invention usually comprises a support having thereon an image-receiving layer. The support may be a transparent film such as poly(ethylene terephthalate) or reflective.

The image-receiving layer may comprise, for example, a polycarbonate, a polyurethane, a polyester, polyvinyl chloride, poly(styrene-co-acrylonitrile), poly(caprolactone) or mixtures thereof.

20 As noted above, the donor elements of the invention are used to form a transfer image. Such a process comprises imagewise-heating a donor element as described above and transferring a fluorescent material image to a receiving element to form the transfer image.

The donor element of the invention may be used in sheet form or in a continuous roll or ribbon. If a continuous roll or ribbon is employed, it may have only the fluorescent europium complex thereon as 25 described above, with or without an image dye, or may have alternating areas of different dyes, such as sublimable magenta and/or yellow and/or cyan and/or black or other dyes.

In a preferred embodiment of the invention, the donor element comprises a poly(ethylene terephthalate) support coated with sequential repeating areas of magenta, yellow, and cyan dye and the fluorescent material as described above, and the above process steps are sequentially performed for each color to 30 obtain a three-color dye transfer image containing a fluorescent image.

If a laser is used to transfer dye from the dye-donor to the receiver, then an absorptive material is used in the dye-donor. Any material that absorbs the laser energy may be used such as carbon black or non-volatile infrared-absorbing dyes or pigments which are well known to those skilled in the art.

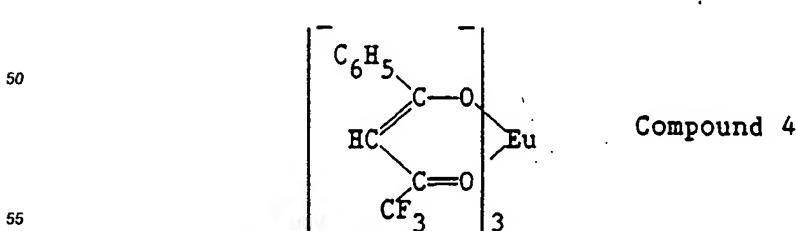
A thermal transfer assemblage of the invention comprises
35 a) a donor element as described above, and
 b) a receiving element as described above,
the receiving element being in a superposed relationship with the donor element so that the fluorescent material layer of the donor element is in contact with the image-receiving layer of the receiving element.

The following examples are provided to illustrate the invention.

40 Preparation 1

Hexacoordinate complexes are obtained by solution reaction of three moles of a diketone ligand and one mole of a soluble europium salt under mildly alkaline conditions.

45 Compound 1 is available commercially from Kodak Laboratory and Research Products. Compound 4 was prepared in the following manner (other 6-coordinate complexes may be prepared in a similar manner):



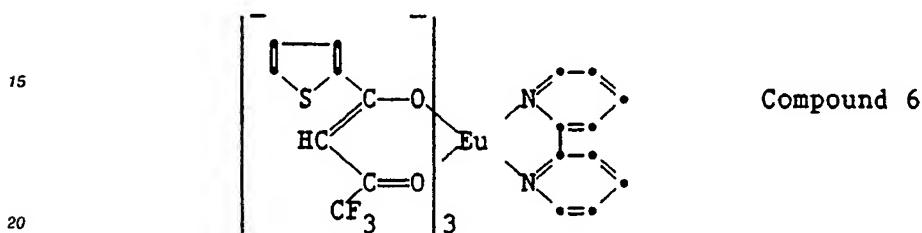
Europium nitrate (450 mg, 1.0 mmole) and benzoyl trifluoroacetone (650 mg, 3.0 mmole) were dissolved in

ethanol (15 ml) and stirred with warming. The solution was adjusted to approximately pH 8.5 with tetramethylammonium hydroxide (25% in methanol) and allowed to stir for 15 min. After this time, the product was aided in precipitating by the addition of water (50 ml), filtered, and dried.

5 Preparation 2

Octacoordinate complexes involving a bidentate auxiliary ligand are obtained by reaction of one mole of a hexacoordinate complex with one mole of the desired neutral ligand.

10 Compound 6 was prepared in the following manner (other 8-coordinate complexes may be prepared in a similar manner):



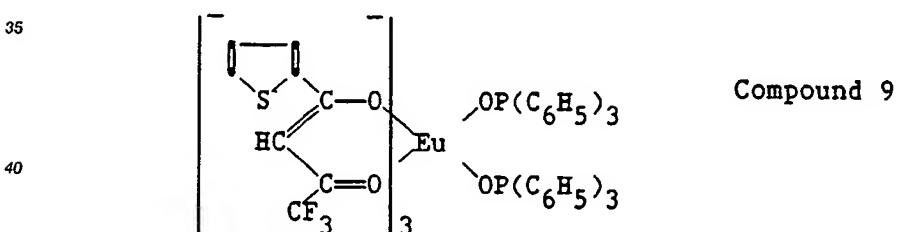
Europium(III) thenoyltrifluoroacetone trihydrate (Compound 1) (860 mg, 1.0 mmole) was dissolved in ethanol (10. ml). 2,2'-Bipyridine (160 mg, 1.0 mmole) dissolved in ethanol (10. ml) was added with stirring.

25 After 15 min, 20 ml of water was added and the resulting precipitate was filtered and dried.

Preparation 3

Octacoordinate complexes involving a monodentate auxiliary ligand are obtained by reaction of one mole of a hexacoordinate complex with two moles of the desired neutral ligand.

30 Compound 9 was prepared in the following manner (other 8-coordinate complexes may be prepared in a similar manner):

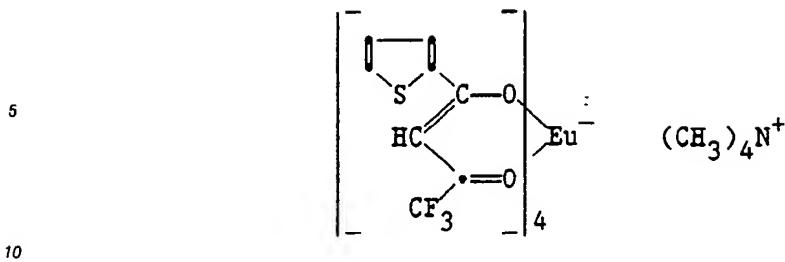


45 Europium(III) thenoyltrifluoroacetone trihydrate (Compound 1) (870 mg, 1.0 mmole) and triphenylphosphine oxide (556 mg, 2.0 mmole) were dissolved in ethanol (8. ml with stirring. After 10 minutes, the resulting precipitate was filtered and dried.

Preparation 4

50 Octacoordinate complexes involving only diketone ligands are obtained by reaction of one mole of a soluble europium salt with four moles of a diketone ligand.

Compound 17 was prepared as follows.



Europium nitrate (450 mg, 1.0 mmole), thenoyltrifluoroacetone (889 mg, 4.0 mmole), and tetramethyl ammonium hydroxide (1.46 g, 25% in methanol, 4.0 mmole) were dissolved in ethanol (20. ml) and heated to boiling. After slowly cooling to room temperature, water (20. ml) was added. The resulting precipitate was filtered and dried.

Example 1

This example shows the fluorescence obtained by thermally transferring 6-coordinate europium complexes from a donor to a receiver.

A donor element was prepared by coating the following layers in the order recited on a 6 μm poly(ethylene terephthalate) support:

- 20 1) a subbing layer of duPont Tyzor TBT[®] titanium tetra-n-butoxide (0.12 g/m²) from 1-butanol; and
- 2) a layer containing the europium fluorescent complex of the diketone ligand as identified above (0.38 g/m²) or control material identified below (0.16 g/m²) in a cellulose acetate butyrate (17% acetyl and 28% butyryl) binder (0.43 g/m² except control at 0.32 g/m²) coated from a cyclopentanone, toluene and methanol solvent mixture.

On the back side of the element was coated:

- 25 1) a subbing layer of duPont Tyzor TBT[®] titanium tetra-n-butoxide (0.12 g/m²) from 1-butanol; and
- 2) a slipping layer of Emralon 329[®] poly-tetrafluoroethylene dry film lubricant (Acheson Colloids) (0.54 g/m²) and S-Nauba 5021[®] Carnauba Wax (Shamrock Technology) (0.003 g/m²) and coated from a n-propyl acetate, toluene, 2-propanol and 1-butanol solvent mixture.

Control Materials

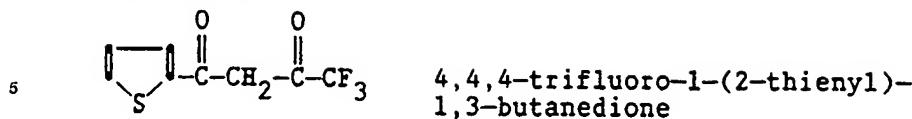
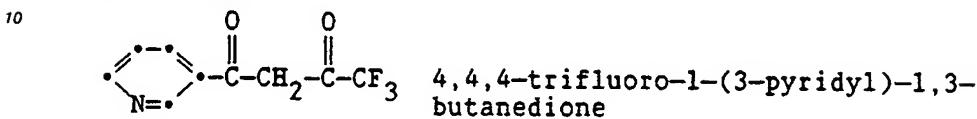
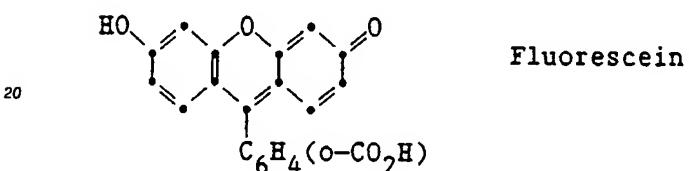
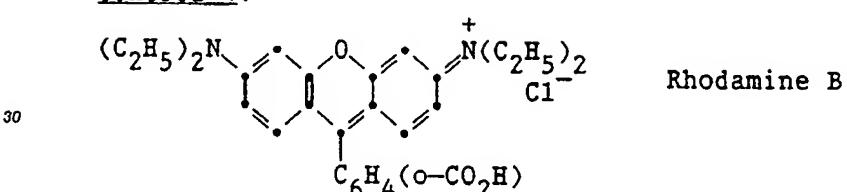
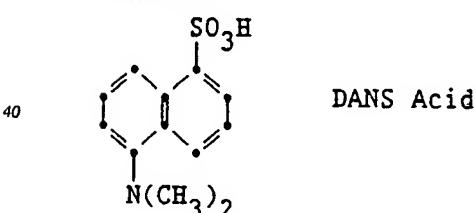
The following materials are available commercially from Kodak Laboratory Products and Chemicals Division:

40

45

50

55

Control 1:Control 2:Control 3:Control 4:Control 5:

45 A receiving element was prepared by coating a solution of Makrolon 5700® (Bayer A.G. Corporation) a bisphenol-A polycarbonate resin (2.9 g/m²) and FC-431® surfactant (3M Corporation) (0.16 g/m²) in a methylene chloride and trichloroethylene solvent mixture on a transparent 175 µm polyethylene terephthalate support subbed with a layer of poly(acrylonitrile-co-vinylidene chloride-co-acrylic acid) (14:79:7 wt ratio) (0.05 g/m²).

50 The fluorescent material layer side of the donor element strip approximately 9 cm x 12 cm in area was placed in contact with the image-receiving layer of the receiver element of the same area. The assemblage was fastened in the jaws of a stepper motor driven pulling device. The assemblage was laid on top of a 14 mm diameter rubber roller and a TDK Thermal Head L-133 (No. 6-2R16-1) and was pressed with a spring at 55 a force of 36 N against the donor element side of the assemblage pushing it against the rubber roller.

The imaging electronics were activated causing the pulling device to draw the assemblage between the printing head and roller at 3.1 mm/sec. Coincidentally, the resistive elements in the thermal print head were pulsed at a per pixel pulse width of 8 msec to generate a maximum density image. The voltage supplied to

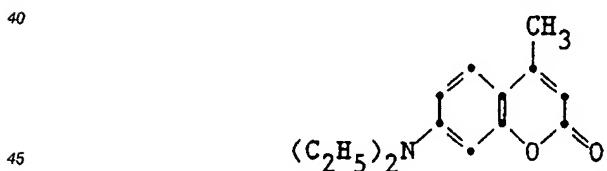
the print head was approximately 25 v representing approximately 1.6 watts/dot (13 mjoules/dot).

The receiving element was separated from the donor element and the relative emission was evaluated with a spectrofluorimeter using a fixed intensity 360 nm excitation beam and measuring the relative area under the emission spectrum from 375 to 700 nm. The following results were obtained (all transferred 5 europium complexes emitted between 610 and 625 nm):

Table 1

	<u>Compound</u>	<u>Relative Emission*</u>	<u>Visual Color</u>
10	None	<1	Not visible
15	Comparison*	100	Blue
20	Compound 1	5	Moderate red
25	Compound 2	5	Moderate red
30	Compound 3	1	Faint red
35	Compound 4	3	Moderate red
40	Compound 5	1	Faint red
45	Control 1	<1	Not visible
50	Control 2	<1	Not visible
55	Control 3	<1	Not visible
60	Control 4	<1	Not visible
65	Control 5	<1	Not visible

* Compared to the following compound, normalized to 100 (emission between 400-500 nm).



This compound is the subject of U.S. Patent 4,876,237.

50 The above results show that the compounds of the invention have a unique red fluorescence and more intense transferred images than the control compounds of the prior art.

Example 2

55 This example is similar to Example 1 but shows the enhanced fluorescence obtained with 8-coordinate europium complexes as compared to the corresponding 6-coordinate complex.

Donor elements were prepared as described in Example 1, using the fluorescent europium complex of the diketone and auxiliary ligand identified above.

Receiving elements were prepared as in Example 1.

The evaluation of fluorescence was done as described in Example 1. The following results were obtained:

5

Table 2

	<u>Compound</u>	<u>Relative Emission*</u>	<u>Visual Color</u>
10	None	<1	Not visible
	Compound 1	5	Moderate red
15	Compound 6	27	Intense red
	Compound 7	32	Intense red
	Compound 9	54	Intense red
20	Compound 10	20	Red
	Compound 17	32	Intense red

25

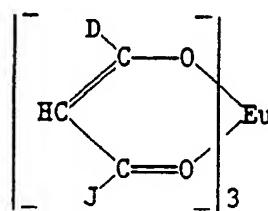
* Compared to the same compound as in Example 1.

compounds of the invention have a unique red fluorescence and more intense transferred images than the control compounds of the prior art.

30 **Claims**

1. A donor element for thermal transfer comprising a support having on one side thereof a fluorescent europium complex dispersed in a polymeric binder.
- 35 2. The element of Claim 1 characterized in that said complex has the formula:

40



45

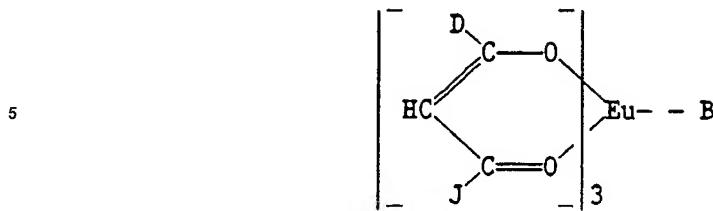
wherein:

D is a substituted or unsubstituted, aromatic, 5- or 6-membered carbocyclic or heterocyclic moiety; and J is -CF₃, -CH₃, -CH₂F or -CHF₂.

50

3. The element of Claim 2 characterized in that D represents phenyl, 2-thienyl, 2-furyl or 3-pyridyl.
4. The element of Claim 1 characterized in that said complex has the formula:

55



wherein:

D is a substituted or unsubstituted, aromatic, 5- or 6-membered carbocyclic or heterocyclic moiety;

J is -CF₃, -CH₃, -CH₂F or -CHF₂; and

B represents at least one monodentate ligand with an electron-donating oxygen or nitrogen atom or at least one bidentate ligand with two electron-donating oxygen, nitrogen or sulfur atoms capable of forming a 5- or 6-membered ring with the europium atom.

20

5. The element of Claim 4 characterized in that B represents tri-n-octylphosphine oxide, pyridine-N-oxide or triphenylphosphine oxide.
6. The element of Claim 4 characterized in that B represents 2,2'-bipyridine, 1,10-phenanthroline, ethylene diamine or 1,2-diaminobutane.
7. The element of Claim 4 characterized in that D represents phenyl, 2-thienyl, 2-furyl or 3-pyridyl.

25

8. The element of Claim 1 characterized in that said donor element comprises sequential repeating areas of magenta, yellow and cyan dye, and said fluorescent complex.
9. A process of forming a transfer image comprising imagewise-heating a donor element comprising a support having on one side thereof a layer comprising a material dispersed in a polymeric binder, and transferring an image to a receiving element to form said transfer image, characterized in that said material is a fluorescent europium complex.

30

10. A thermal transfer assemblage comprising:
 - a) a donor element comprising a support having on one side thereof a layer comprising a material dispersed in a polymeric binder, and
 - b) a receiving element comprising a support having thereon an image-receiving layer, said receiving element being in a superposed relationship with said donor element so that said material layer is in contact with said image-receiving layer, characterized in that said material is a fluorescent europium complex.

45

50

55